

Air Emission Inventories in North America: A Critical Assessment

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ABSTRACT

Although emission inventories are the foundation of air quality management and have supported substantial improvements in North American air quality, they have a number of shortcomings that can potentially lead to ineffective air quality management strategies. Major reductions in the largest emissions sources have made accurate inventories of previously minor sources much more important to the understanding and improvement of local air quality. Changes in manufacturing processes, industry types, vehicle technologies, and metropolitan infrastructure are occurring at an increasingly rapid pace, emphasizing the importance of inventories that reflect current conditions. New technologies for measuring source emissions and ambient pollutant concentrations, both at the point of emissions and from remote platforms, are providing novel approaches to collecting data for inventory developers. Advances in information technologies are allowing data to be shared more quickly, more easily, and processed and compared in novel ways that can speed the development of emission inventories. Approaches to improving quantitative measures of inventory uncertainty allow air quality management decisions to take into account the uncertainties associated with emissions estimates, providing more accurate projections of how well alternative strategies may work. This paper discusses applications of these technologies and techniques to improve the accuracy, timeliness, and completeness of emission inventories across North America, and outlines a series of eight recommendations aimed at inventory developers and air quality management decision makers to improve emission inventories and enable them to support effective air quality management decisions for the foreseeable future.

INTRODUCTION

Air quality management (AQM) in North America focuses on ensuring that concentrations of compounds in the ambient air are below the levels that are considered harmful to human health or the environment. Strategies developed to achieve these standards are based upon reduction of emissions from specific source classes. The effectiveness of this approach depends upon an accurate understanding of the relative contributions of the sources to ambient atmospheric pollution.

An adequate knowledge of emissions sources and associated fluxes, both before and after emission controls are adopted, has long been recognized as a requirement for designing cost effective air pollution control strategies.¹ Emission inventories are designed to systematically quantify the temporal and spatial distributions of the fluxes of primary pollutants and secondary pollutant precursors emitted by significant sources. This places emission inventories at the foundation of today's air quality management strategies, and significant errors in inventories can, therefore, lead to the adoption of strategies that protect human health and the environment less effectively than possible. Emission inventory errors can be enormously expensive, by requiring installation and operation of air pollution controls beyond the minimum needed, and by failing to effectively reduce adverse health and environmental damage.

The purpose of this paper is to identify the status of current emission inventory practices, point out the general strengths and weaknesses of existing inventories, and to suggest possible directions for improving future inventories. The suggested directions are based upon the recommendations developed as part of the recent assessment of emission inventories by NARSTO.²

BACKGROUND

Several recent reports have recognized the importance of emission inventories and the challenges that must be overcome to ensure that inventories are able to provide the quality of information needed to support sound AQM decisions. In the latest of these reports, the National Research Council (NRC) noted in 2004 that, "The first step in

developing an emission-control strategy for a criteria pollutant is to develop an inventory of pollutant emissions that lists all sources of the pollutant or its precursor and the rate at which each source emits the pollutant to the atmosphere.”³

In response to the NRC’s recommendations, the Clean Air Act Advisory Committee (CAAAC), an advisory committee to the U.S. Environmental Protection Agency (EPA), created an Air Quality Management Work Group to develop a plan to address the report’s recommendations. The AQM Work Group agreed with the need to strengthen emission inventories to ensure adequate support for AQM strategy development, and concluded that, “A strong national effort is needed to improve emission estimation methods for major source categories, especially for sources that are poorly characterized or whose emissions estimates are uncertain.”⁴ In other scientific reviews conducted by the NRC^{5, 6} and NARSTO,^{7, 8} emission inventories have consistently been seen as needing improvement to enable them to continue to be of use in the development of effective AQM strategies.

Given the consistent call for improvements in inventories of air pollutant emissions by these diverse expert panels, it would be easy to conclude that air pollutant emission inventories are severely flawed and are of little effective use to air quality managers. However, measurements show that U.S. emissions of the pollutants addressed in the original Clean Air Act have decreased, in some cases enormously, over the past 20 years. Ambient concentrations of those pollutants have also decreased significantly, even though economic and personal activities responsible for those pollutant emissions have increased considerably over that period.⁹ This would suggest that air quality managers have had a good understanding of what emissions sources to control in order to improve air quality.

Each of the expert panels cited above examined specific applications of emission inventories and the ability of emission inventories to support future air quality management decision making. Although national emission inventories are currently capable of estimating the average annual emissions on a national scale, those same

inventories have shortcomings when used in other contexts, such as estimates of daily emissions in a local area.

Most air quality management goals have focused on emissions from major, and relatively well characterized, source categories. As recently implemented regulatory programs take effect, however, emissions from these sources will decline substantially. The remaining emissions will be more evenly distributed over source categories that are more difficult to measure or model. A key example of this changing context is the recent promulgation in the U.S. of the Clean Air Interstate Rule (CAIR)¹⁰ and the rules to reduce emissions from on- and off-road diesel vehicles.^{11, 12} Together these rules address emissions from the largest source categories of nitrogen oxides (NO_x) and SO₂, mobile sources and coal-fired electric generating units, respectively. As these rules begin to take effect, these sources will no longer be as dominant in the total U.S. emissions of NO_x and SO₂ (see Figure 1), making other source categories relatively much more important.¹³ In this and similar situations, errors in emission estimates from smaller individual sources will have greater consequences than were previously the case. These consequences could range from wrongly identifying a pollutant that should be controlled to overlooking source categories whose control could result in more cost-effective emission reductions.

In addition to this loss of dominating sources, future air quality management strategies are beginning to consider all emissions into an airshed, including hazardous air pollutants (or air toxics) that have previously been considered separately from the criteria pollutants. This approach is consistent with one of the NRC Air Quality Management recommendations to evaluate the entire load of pollutants entering and emitted within an airshed, rather than managing them individually.³ To address this recommendation, inventories of all pollutants would need to be at least compatible with one another, and ideally integrated into a single coherent inventory.

The changing context in which emission inventories are being used, a greater need to understand the limitations associated with emission inventory data, and a higher expectation for rapid and flexible data availability are placing tremendous pressure on the

166 developers of emission inventories to provide accurate, timely, accessible, and flexible
167 emission inventory databases. To address these issues, NARSTO recently released an
168 assessment of emission inventories across North America, which included eight
169 recommendations for improvements.² Although the focus of this assessment was on
170 national emission inventories, the recommendations are applicable to international,
171 regional, state and provincial, and local inventories as well. It must also be noted that the
172 recommendations were developed with a North American, rather than strictly a U.S.,
173 perspective. Even though the NARSTO assessment identified differences in emission
174 inventory development and needs across Canada, Mexico, and the U.S., the majority of
175 issues and fundamental needs are common to all three countries. Much of the following
176 discussion is drawn from the NARSTO emission inventory assessment, with the same
177 focus on North American inventories.

178 179 **EMISSION INVENTORY EVOLUTION AND STRUCTURE**

180 Most early emission inventories were developed to help address air quality problems
181 around specific major metropolitan areas,¹⁴⁻¹⁶ as recognition grew during the 1950s and
182 into the 1960s that air pollution was a significant public health problem.¹⁷ To more
183 accurately estimate emissions in these areas with the technology available at that time,
184 factors were developed that related emissions to industrial and other activities. These
185 emission and activity factors allowed air quality managers to estimate how changes in
186 activity levels or technologies impacted total emissions without requiring measurements
187 at every facility, and this approach has formed the foundation of modern emission
188 inventories. Although early emission inventory developers confronted many problems
189 similar to those faced by current efforts, there are also some significant differences. Our
190 sampling methods, for instance, have fortunately advanced considerably since Rossano
191 and Schell stated that, “Observing the effluents where possible, and even smelling or
192 feeling them may provide useful information.”¹⁸

193
194 In their most simple form, emission inventories are developed using emission factors
195 (*EFs*) and associated activity (*A*) information. Emission factors are the mass of pollutant
196 emissions released per unit of the associated process variable. Activities are the related

process variable, such as mass of fuel consumed or output produced. The emissions (E) are then calculated as:

$$EF \times A \times [1 - (ER/100)] = E \quad (1)$$

where ER is the emission reduction (in percent) associated with use of a pollution control system.

In lieu of using an emission reduction factor, a different EF can be used, particularly if the pollution control approach involves a modification of the process. The $EF \times A$ structure assumes that the emission factor is independent of the activity factor. Where the activity is a process variable, e.g., load or throughput, this assumption may not be accurate, as it is possible for the emission factor to vary as the process varies. On the other hand, when the activity measure is the population of similar sources such as the number of dry cleaners, the assumption of independence is reasonable.

Over a large number of sources and a long period of time, variations in the emission rate can be expected to even out so that the emission factor adequately reflects the average emission rate across the activity range; this allows emission factors to be used as the basis for developing national annual emission inventories. The U.S. EPA compiles a database of emission factors for a wide variety of source types and a range of pollutants in its “AP-42” document. The stationary source volume of AP-42 is currently in its fifth edition, and has 15 chapters covering sources as diverse as external combustion sources (such as boilers and process heaters), storage tanks, and ordnance detonation.¹⁹

In spite of the long history of the emission factor approach, it does have significant shortcomings. As far back as the 1950s, issues such as temporal and spatial allocation of emissions or nonlinear relationships between activity and emission levels were recognized.²⁰ Even fifty years later, these complexities continue to present problems for emission inventory developers. To address some of these issues, emission models have been developed to more accurately estimate emissions from sources with complex

operating characteristics that cannot be accurately represented by the simple relationship in Eq. (1). The more well-known of these models have been developed for estimating mobile source emissions. Although early mobile source inventories used an average emission factor and fuel consumption data to estimate emissions, current emission models account for changes in fuel type, fuel evaporation, engine deterioration, operation of air conditioning, and engine startup in addition to the variation in emissions due to differences in vehicle design.

Over the years, emission models have grown in scope and complexity. Models for on-road and non-road mobile sources have been developed by the U.S. EPA and others.²¹⁻²⁴ Other emission models have been developed to estimate emissions from vegetation and soil.²⁵ Pollutant-specific models have also been developed, such as the Carnegie-Mellon University ammonia (NH₃) emissions model, which estimates NH₃ emissions from animal feeding operations, wastewater treatment, and mobile sources, as well as from natural processes.²⁶

Figure 2 illustrates how these different components — emission factors, activity factors, and emission models — combine to create a complete inventory.²⁷ For stationary point sources, one can determine emissions using the appropriate emission factors combined with the facility-specific activity factors. Emissions for non-point sources are estimated using the EPA MOBILE, NONROAD, and Biogenic Emission Inventory System (BEIS) models, among others. These annual emission estimates are then combined to determine the total estimated emissions. However, modern inventories involve many additional elements to ensure that they are as relevant as possible to current air quality management needs.

CURRENT INVENTORIES

The growth of emission models is a reflection of the increasing scope and complexity of emission inventory needs, and of emission inventories themselves. The general structure of the current national emission inventories (NEIs) was derived from the early metropolitan-area inventories beginning in the 1970s in the U.S. and Canada.²⁸ The

1985 National Acid Precipitation Assessment Program (NAPAP) emission inventory was EPA's first attempt to produce a highly quality-assured national inventory for use by policy makers, modelers, human and ecological effects researchers, and industry, and set the stage for today's NEIs.²⁹ These inventories (and this paper) focus on criteria pollutant emissions, and typically include CO, NH₃, NO_x or NO₂, PM (including specific size fractions such as PM₁₀ and PM_{2.5}), SO₂, and VOCs.^{2, 30}

The NEIs in turn have provided the structure, and often the data, for emission inventories at the local, state and provincial, and regional levels. These inventories are developed to address more specific air quality management issues, such as regional haze, implementation of local air quality management plans, or cross-border air quality problems.³¹⁻³⁴ Specialty emission inventories, such as those for dioxins and dioxin-like compounds,³⁵ mercury,^{2, 36} and black carbon³⁷ are also needed to address emissions of specific pollutants. These regional and specialty inventories generally follow the NEI approach that relies on emission and activity factors and emission models as a guide to determine what information, and in what format, is needed. Each of the variations on the NEI challenges the ability of the basic inventory structure to meet the specific inventory needs. Probably the greatest challenge modern inventories face, however, is their use as the basic source of emissions data for air quality models.

The need to design and implement air quality management plans to reduce ambient concentrations of ozone has driven the development of increasingly complex models of atmospheric transport and photochemical reactions, and the atmospheric formation of secondary PM is continuing to push for even more complexity. Because many of the ozone formation processes occur in the atmosphere on time scales of hours or less, and since the chemistry included in these models is necessarily complex, these models require increasingly detailed information on the location, time, and chemical speciation of emission fluxes of the major ozone precursors, NO_x and VOCs.³⁸ Because emissions data are generally available as annual averages (except for some major source categories such as electric generating units), annual emission inventory data are fed into processing models (emission processors) that allocate emissions temporally (and spatially for area

sources) to simulate the actual hourly, daily, and seasonal changes in emissions. Such simulations must account not only for these temporal emission changes, but also for changes due to changes in meteorological conditions. Processes such as space heating and cooling, evaporative and biogenic emissions, and even traffic patterns change as meteorology changes. These effects add further complexity that must be accounted for when developing emission inventories.

Emissions processors operate upon the base inventory data to provide model-ready detailed emissions, as illustrated in Figure 3.³⁹ Processors such as the Sparse Matrix Operator Kernel Emissions (SMOKE) model,⁴⁰ the Emissions Processing System (EPS), and the Emissions Modeling System (EMS) generate input files for the atmospheric models. These processors typically provide hourly emissions over the course of a week, allocated to grid cells over the model domain, and with the speciation required by the chemistry model being used. The detailed chemical species are typically contained in a separate database that lists speciation profiles for a range of chemical compounds usually emitted by a given source type. A database that has been widely used is EPA's SPECIATE, which contains speciation profiles for a wide range of source categories.⁴¹

As this introduction has illustrated, the scope of the applications of emission inventories is thus extremely broad, and can range from hours to decades on a temporal scale, and from neighborhood to global on a spatial scale. These widely differing scales and purposes can result in significant mismatches between the emission inventory data and the needs for those data, given the temporal or spatial scales or degree of speciation available. These mismatches act to highlight the shortcomings of existing emission inventories, and these will be discussed in more detail below. But although emission inventories are not yet ideal, considerable work has been done over the past 30 years to strengthen their ability to provide critical information for developing successful air quality management strategies.

EMISSION INVENTORY STRENGTHS

Current emission inventories can, in general, be used to compare the relative significance of different source categories. Major insights can be drawn from the current U.S. NEI – the largest fraction of NO_x and SO₂ emissions and a considerable portion of VOC emissions are from stationary sources; mobile sources are the largest contributor to total CO emissions and a considerable contributor to total NO_x and VOC emissions; and biogenic sources contribute the largest portion of total VOC emissions.^{9, 42} On a national annual emissions basis, these insights enjoy a high degree of confidence.² Similarly, air quality managers in Canada and Mexico are able to identify the key sources of concern based on information derived from their respective national inventories.

Existing emission inventories provide insight into air-quality trends over time and overall pollution control efficiency. Comparison of current emission inventories to those from previous years provides the basis for estimating emission trends over time. Such comparisons can give an indication of efficiency of particular control strategies, particularly those that are national in scope, such as reductions in SO₂ and NO_x from large point sources in the United States associated with acid rain provisions of the Clean Air Act Amendments of 1990.

Air quality management strategies can be developed based upon current emission inventories. In addition, priorities for air quality improvements can be set based upon the knowledge of emissions contributed from different source categories.⁴³ With limited resources, such prioritization focuses efforts on those sources with the greatest potential to reduce emissions. For example, in urban areas facing ozone problems, the relative importance of NO_x versus VOC control can be assessed taking into account both urban scale and regional geographic scales, and the key source categories that should be the focus of control efforts can be broadly prioritized.

Many of the tasks required to develop current emission inventories are more efficiently completed using existing emission models and tools. Tools such as MOBILE, TANKS (for estimating VOC and HAP emissions from organic liquid storage tanks) or the Emission and Dispersion Modeling System (EDMS) (for estimating emissions from

aircraft and airport operations) help automate many of the tasks involved in developing emission inventories, thereby reducing the cost of inventory development. A further benefit is that the use of many of these tools leads to the standardization of emission inventory development efforts, thereby increasing their compatibility across emission inventories.²

INVENTORY NEEDS AND SHORTCOMINGS

In an ideal situation, the information reported in the different emission inventories would all be based upon a single set of basic data that could then be compiled to provide the information needed at a particular time for a particular application. The basic data would be updated frequently as new information was received, and would include the appropriate metadata describing the information source, data collection methods used, limitations (including variability, uncertainty, and applicability), and other descriptors. Inventories would be compiled upon request, and updates would be available within days, if not hours, of new information being submitted.

The NRC's Air Quality Management panel emphasizes such a need for compatible inventory data, given the stress on managing airsheds as a single entity rather than managing individual pollutants. Although the latter approach is driven by the regulatory structures created (in the U.S.) by the Clean Air Act, the NRC points out that the appropriate scientific structure is to evaluate emissions and air quality management on a scientifically-based foundation of a single airshed that will be affected by each AQM strategy regardless of its regulatory basis.³ Thus, the NRC's recommendations apply equally to emission inventories across North America.

In such a situation, different inventories would be compatible and comparable, and near-real-time. Users would be able to understand how certain the data were, and would have the ability to evaluate how uncertainties propagated through analyses that used the inventory data. Air quality managers and policy makers would have the latest information on emissions and a measure of how the data would be likely to change based upon the certainty and variability information.

382
383 Unfortunately, we are far from this ideal situation. There can be considerable uncertainty
384 in the data reported in emission inventories, for a number of reasons.^{44, 45} The
385 shortcomings of current emission inventories are closely related to these sources of
386 uncertainty. For instance, many, if not most, source emission estimates are based on a
387 small number of measurements that do not adequately represent the full range of process
388 designs and operational practices. This limits the accuracy and increases, sometimes
389 significantly, the uncertainty of the estimate.^{3, 6} Inventories that rely on emission factors
390 typically do not account for changes in emissions during startup and shutdown, or often
391 during significant operational changes. Although emissions models can estimate these
392 emissions, they are not often included, particularly for stationary sources.² Area source
393 emissions are often spatially allocated at the county level, requiring estimates of spatial
394 allocation factors to be used if the inventory data are to be used for air quality modeling.
395 Similarly, emission inventories are usually based upon annual average values, which may
396 be far from actual values over the hour-long time frames of interest to air quality
397 modelers.

398
399 The emission inventories developed at the county, state or provincial, and national levels
400 are often inconsistent. Data from different agencies may be developed using different
401 estimating procedures, several similar emissions source types may be consolidated, and
402 information regarding data use limitations may not be carried through as data are
403 combined to create inventories that cover a larger geographical range. Conversely, data
404 that are adequate at a national level can become much more uncertain when
405 disaggregated to estimate emissions at a state or local level due to differences in the
406 source mix, operating conditions, and other factors.² Mechanisms are needed to ensure
407 that the aggregation or disaggregation process is appropriate and consistent within and
408 across different countries and, in some cases, across agencies within a country.

409 Uncertainties are greater for some sources and pollutants than for others. Emissions of
410 NO_x and SO₂ from electric utility generating units (EGUs) in the U.S., for instance, are
411 measured using continuous emissions monitors, and the data are reported to the U.S. EPA
412 each quarter. These data are accurate and well-resolved spatially and temporally, but

data on other pollutants and other sources are much more difficult to obtain. In particular, emission inventories of hazardous air pollutants are more uncertain than criteria pollutants, and emissions from natural, area, and mobile sources are more uncertain than those from large stationary point sources. Emissions from anthropogenic sources are much more well-characterized than emissions from natural sources, and, hence, the emission estimates for natural sources are often much more uncertain.⁴² A relative comparison of the qualitative confidence levels for national inventories of key gaseous pollutants is shown in Table 1.^{2, 7}

Uncertainties also arise because of the time intervals between updating and reporting of emission inventories, and between sampling and reporting of emissions data. As the period between data collection and data reporting lengthens, the uncertainty of the reported data to represent actual emissions increases. The U.S. NEI is updated every three years, and thus may not provide timely and updated emission information for air-quality management decisions. In addition, many of the emission factors and speciation profiles are based on measurements that are over a decade old, resulting in questionable applicability of the measurements.² Efforts are being made to reduce these intervals, notably in Canada, which has as a goal annual updates to their emission inventories.

Typically, the uncertainties associated with the inventory data are not reported, and often, the information needed to quantify the uncertainties is not collected. Quality assurance and quality control procedures are not strictly applied in most emission models or during inventory development, resulting in unknown levels of data certainty. For most emission inventories, documentation of the key assumptions and data sources used during the emission inventory development is inadequate. Although the AP-42 emission factors compilation provides some guidance on the uncertainty of emission factor data, the quality ratings are not quantitative.¹⁹ It should be noted that EPA is in the process of revamping the emission factor program. One goal of this program is to provide quantitative uncertainties and guidance on the use of factors.

FUTURE DIRECTIONS AND TRENDS IN INVENTORY DEVELOPMENT

Many new tools and techniques have been evaluated and applied to reduce the uncertainties associated with emission inventories. Most of these techniques are associated with emission measurements, but a number of developments in information technology (IT) and computational models are improving inventories through better information processing and expanded capabilities to evaluate inventories' accuracy. In many cases, combining innovative measurement and computational techniques can provide new approaches for improving emission inventories.

Remote Sensing

Many of the new emission measurement technologies rely on remote sensing rather than on conventional "probe-in-stack" approaches. Ground-based remote sensing methods that rely on absorption spectroscopy include non-dispersive infrared (NDIR) techniques, Fourier transform infrared (FTIR) methods, differential optical absorption spectroscopy (DOAS), and tunable infrared laser differential absorption spectroscopy (TILDAS).² FTIR has been used to measure NH₃ emissions from animal feeding operations,^{46, 47} a range of organic compounds from combustion processes,⁴⁸ and emissions from natural gas flares.⁴⁹ DOAS measurements have been combined with dispersion modeling to estimate emissions from gasoline service stations and tanker filling operations.⁵⁰ Other ground-based remote sensing methods rely on fluorescence or Raman spectroscopy or light detection and ranging (LIDAR) techniques.⁵¹

Ground-based remote sensing has been especially useful in evaluating emissions from mobile sources. The most common approach has been to conduct measurements across roadways and intercept the exhaust plumes from vehicles moving across the measurement path. NDIR techniques were used in some of the earliest studies to measure CO and CO₂, and more recently emissions of HC and NO_x.⁵²⁻⁵⁶ More advanced TILDAS applications have been developed to measure a wider range of species, including several nitrogen and organic compounds.⁵⁷ Still in the development stage are dispersive infrared (IR) techniques, use of LIDAR, and instrumentation that can measure exhaust PM.⁵⁸⁻⁶⁰ Future directions for this technique include the use of remote sensing to quantify air

toxics, PM and PM precursors, and greenhouse gases.^{61, 62}

Satellite and Aircraft-Based Sensors

Numerous remote sensing applications are based on satellites or aircraft. Satellite measurements are currently being used to visually identify the location and intensity of large-area fires, ship plume tracks, major industrial plumes, and incidents of dust storms or regional haze.^{2, 63} Measurements of surface properties such as ground cover and temperature may be the most effective current use of satellite data, but progress has been made in using satellite data to infer pollutant column concentrations or densities and continental-scale emissions. Even though the visual images collected by satellites have aided in identifying areas of smoke and haze emissions and transport, these data are limited in capability to provide quantitative data. Numerous technical challenges to collection of quantifiable pollutant concentration data remain, including compensating for variations in the air-chemistry matrix, aerosol burden, cloud cover, surface albedo, and temperature, as well as dealing with masking effects of the stratospheric overburden, which can be dominant. Current technology does not enable effective satellite measurements to be made beneath cloud cover.²

Even so, satellites' ability to cover large, spatial domains is a major advantage over other approaches. Combining satellite measurements with data from other sources, including air quality and atmospheric behavior models, can provide valuable information in instances where no other measurements are available. Estimates of emissions over a large geographical area, of the relative amounts of natural and anthropogenic emissions, and of emissions where little or no quantitative data are available are well-suited for evaluation by satellite-based methods. Perhaps the most effective use of satellite data has been as an independent source of data to which inventory results can be compared to identify inventory gaps and shortcomings. This technique has been applied to biogenic emissions of isoprene, global NO_x emissions, and CO emissions from wildfires.⁶⁴⁻⁶⁶

Aircraft remote sensing is most often used to measure the flux of pollutants that pass through a vertical plane intersecting the pollutant plume that is being observed. DOAS,

IR spectroscopy, or lidar are the typical techniques used in aircraft remote sensing applications.^{67, 68} Current aircraft remote sensing is largely limited to NO_x and SO₂ fluxes using DOAS and organic species from biomass burning using IR spectroscopy. Even so, aircraft measurements of this type are particularly useful in determining pollutant fluxes from spatially-extended area sources, such as forest fires and total emissions from an urban area. There is potential for DOAS techniques to be extended to some VOC species including formaldehyde, alkenes and aromatics and for lidar instrumentation to be applied to ozone and PM.²

Mobile Source Emissions Inventories

Mobile sources present numerous and significant challenges to inventory developers.⁶ Not only are emissions from mobile sources a major contributor to ambient air pollution, there is also enormous variability in the many parameters that influence mobile source emissions in terms of mass and composition. Measuring and estimating those emissions have, therefore, been a major focus of emission inventory development from the outset. Mobile source emissions vary with time of day, week, or year; across and within individual sources; and across and within metropolitan areas. Because we are only able to directly measure emissions from an extremely small fraction of the total number of mobile sources, significant efforts are required to extrapolate those measurements to fleet-wide distributions of emissions. Emission models, remote sensing, and statistical methods are a few of the approaches that are used to more accurately estimate emissions from mobile source categories. Because of the importance and difficulties associated with mobile source emissions, we will devote a greater amount of attention to this sector.

Mobile source emission inventories are primarily developed using emission models that calculate emissions across a designated area, usually on an urban scale. The most widely used model is EPA's MOBILE model (and its derivatives in Canada and Mexico), which is in its sixth major revision.^{21, 22} The MOBILE model relies on emissions data from dynamometer testing over standard operating cycles, as well as local characteristics such as fuel composition, climate, and fleet composition. For example, these local characteristics are considerably different for Mexico than for Canada, and the basic

structure of the MOBILE model allows inventory developers to account for such differences.

EPA's NONROAD model uses similar techniques to estimate emissions from non-road vehicles such as construction and agricultural equipment, railroad locomotives, marine vessels, recreational vehicles, and small engines such as those used in lawnmowers and leaf blowers.²³ The resulting emissions estimates are idealized to the extent that the standard driving cycles and fleet characteristics do not fully represent real world conditions.⁶⁹ Next-generation models, such as EPA's Motor Vehicle Emissions Simulator (MOVES) model, are designed to enable more complex estimates of mobile source emissions by using emission rates based on operational modes that change with location (a "modal" approach).⁷⁰ This allows the model to more accurately estimate the time and location of emissions from operating conditions such as cold starts, extended idling, and heavy acceleration.

Modal emissions models such as MOVES will require more detailed measurements under real world conditions. There are several approaches to measuring emissions during actual operations, including remote sensing, tunnel studies, mobile laboratories and chase vehicles, and portable emission measurement systems (PEMS) and on-board sensors.

Remote sensing is generally able to measure the frequency and impact of high-emitting vehicles, and to evaluate the effectiveness of AQM approaches such as the use of oxygenated fuels and inspection and maintenance (I/M) programs.^{57, 71-75} Extensive deployment of remote sensing equipment can also provide near real-time data on roadway emissions that can be used to maintain current emission inventories and provide better data on spatial and temporal emissions distributions. Although remote sensing is usually used with passenger vehicles, the technique has also been extended to heavy-duty diesel trucks and off-road vehicles.⁷⁶⁻⁷⁸ Remote sensing data do have limitations; in particular they are measurements at a limited number of locations and therefore may not include emissions over the full range of operating conditions, such as cold starts.⁷⁹

Tunnel studies measure pollutant concentrations at the entrance to roadway tunnels and at the outlet of tunnel exhaust air systems as a means to estimate mobile source emissions. Combined with counts of vehicle number and type passing through the tunnel, these measurements can be used to develop fleet-level emission rate distributions.^{69, 80-83} Although these measurements are limited to the particular mix of vehicles under particular conditions, tunnel studies provide information on the accuracy of mobile source emission model predictions for aggregate emissions and can identify discrepancies with the predicted mixture of emitted pollutants.⁶⁹

Mobile source emissions can also be characterized using mobile laboratories and chase vehicles, which measure emissions from other vehicles during on-highway operation in normal day-to-day traffic conditions. Both of these approaches rely upon fast-response instrumentation, usually mounted on truck beds or in trailers. The instrumentation can be as simple as a single monitor for gaseous pollutants or as complex as a full suite of sampling equipment for gases, particles, and operating parameters. For instance, the mobile laboratory described in Kolb et al. is equipped to quantify exhaust emissions of gaseous CO, NO, NO₂, HONO, NH₃, H₂CO, CH₃CHO, CH₃OH, benzene, toluene, C₂-substituted benzenes, and SO₂, as well as a range of PM properties, including number density, size distribution, and mass loadings of SO₄⁼, NO₃⁻, NH₄⁺, OC species, and PAHs.⁸⁴

Mobile laboratories collect data by measuring the elevated pollutant concentrations along the roadway without resolving emissions from specific vehicles. These studies are analogous to tunnel studies with regard to the fleet-level evaluation of emissions.⁸⁴⁻⁸⁶ A number of groups have focused on characterizing on-road exhaust emissions of PM, with some placing particular emphasis on concentration and properties of ultrafine particles (e.g., those with aerodynamic diameters smaller than 100 nm).⁸⁷⁻⁸⁹ CO₂ measurements are often used as markers to which pollutant measurements are correlated to distinguish between background and roadway emissions, and also allow estimates of emissions per unit fuel consumption.

Chase vehicles follow specific vehicles, either cooperating or non-cooperating, and sample the target vehicle's individual exhaust plume. The chase laboratory must shadow the target vehicle and must contain fast response (~1-s) sensors measuring CO₂ and the target pollution of interest. If the target vehicle is cooperating with the chase vehicle, operating parameters can be obtained from sensors on board the target vehicle. Otherwise, speed and acceleration must be inferred from the chase vehicle's speed and acceleration, in combination with range-finding measurements if available. Chase vehicles have been used to characterize emissions from light- and heavy-duty vehicles, both diesel and gasoline.^{84, 90-93} Chase vehicle measurements appear to be comparable to cross-road remote-sensing data, with the advantages that a wider range of operating conditions can be sampled and, in many cases, many more exhaust species and properties can be measured.

Collection of emissions data during actual operation can also be achieved by using instrumentation that is installed and operated directly on the vehicles of interest. Particularly for off-road vehicles, this can provide a means to measure emissions under real-world conditions without significant modification of the vehicle or removal of the engine. The two major approaches to on-vehicle measurements are portable emission measurement systems (PEMS) and on-board diagnostic (OBD) sensors.

PEMS require analytical instruments that are sufficiently compact to be mounted on or inside the vehicle and robust enough to withstand the temperature and vibrations experienced during actual operating conditions. The U.S. EPA has led the development of PEMS, with their first-generation system called the Real-Time On-Road Vehicle Exhaust Gas Modular Flowmeter and Emissions Reporting System, better known as ROVER.⁹⁴ ROVER and its follow-on system, the Simple Portable Onboard Test or SPOT (designed specifically for off-road applications), have the dual goals of providing test beds for improving PEMS technology and encouraging private sector development of similar systems. PEMS are currently being deployed in support of field test programs to evaluate compliance and evaluate in-use vehicle emissions, with particular emphasis on identifying high emitting vehicles.⁹⁵

629

630 Contemporary vehicles include a range of on-board diagnostics, including exhaust-gas
631 oxygen and temperature sensors, and engine-load and fuel-consumption monitors. By
632 installing wireless communications devices to transmit the signals from these sensors,
633 OBDs can provide real-time data that can be used to estimate CO, NO_x and VOC
634 emissions from target vehicles. Although some testing has been conducted using this
635 approach, comparison with other measurement methods has not yet been done.⁹⁶
636 However, OBDs can be used to identify vehicles that are malfunctioning and likely
637 emitting at higher than designed rates. Additional work is needed to compare the ability
638 of OBDs to estimate emissions with PEMS and remote sensing techniques.

639

640 Further development of sensor technologies is leading toward “microsensors” that require
641 low maintenance, low operating power, and can survive in high-temperature, chemically
642 reactive post-combustion environments. The Argonne National Laboratory has been
643 awarded several patents for “smart” microsensors developed from ceramic-metallic
644 (cermet) materials, and which use neural network signal processing to relate electrical
645 signals to gas concentrations.⁹⁷ Tests have been conducted to measure CO₂ and O₂, and
646 the sensors may be configured and “trained” to detect other pollutants such as VOCs. The
647 potential for low manufacturing costs (\$0.25 per sensor) make it possible to consider
648 microsensors for a range of applications beyond mobile sources, including as continuous
649 emission monitoring systems (CEMS) for small stationary sources for which monitoring
650 costs are currently prohibitive.

651

652 **DATA UNCERTAINTY, VERIFICATION, AND QUALITY ASSURANCE**

653 Given the critical roles that emission inventories play in the development of AQM
654 strategies, there is a need to better understand the level of uncertainty associated with
655 inventory data. As noted in the background discussion, as emissions become less
656 strongly dominated by a single source category, the need to quantify the uncertainties
657 associated with emissions data becomes more important to ensure the development of
658 effective AQM strategies. This need has been identified by several review panels, in
659 particular the NRC panel on Air Quality Management in the United States.³

The need to understand uncertainty is driven by a number of factors, most importantly whether the uncertainty in the inventory data is significant enough to impact the effectiveness of proposed AQM strategies, or whether reported differences between alternative strategies are meaningful.² Uncertainties in inventory data are a consequence of uncertainties, variabilities, data entry and other errors, and assumptions in emission measurements, activity data, and emission models. These uncertainties further propagate through air quality models and projections of future emissions, both of which have their own additional uncertainties.^{42, 69} The impact of emission data uncertainties can therefore be very complex and difficult to isolate. Nevertheless, there are approaches that can be used to quantify uncertainties in emissions data and in its subsequent use. Numerous examples of uncertainty analyses have been reported for evaluations of emissions of NO_x, VOCs, greenhouse gases, selected hazardous air pollutants, and biogenic emissions.^{43, 98-104} Among the source categories examined in these evaluations are highway vehicles, nonroad vehicles, electric generating units, and biogenic sources.

The impacts on air quality model results due to uncertainties in emission inventory inputs have been evaluated for several cases, particularly for ozone modeling.¹⁰⁵⁻¹⁰⁹ In some studies, the key variables that had the strongest influence on air quality predictions were identified, providing guidance to air quality managers regarding the information of most importance and where additional efforts should be made to improve emission data quality.

Beyond the approach of using direct measurement data to evaluate the uncertainties associated with emission inventory data, the use of “top-down” evaluations can also provide information regarding emission inventory uncertainty. Top-down evaluations use data sources independent of those used to develop the emission inventory, but still closely related to emissions, as the basis for comparison to inventory data. If chosen correctly, these independent data can provide critical tests of an inventory’s accuracy.

There are several techniques that can be used in top-down emission inventory evaluations. A common approach is to compare ambient measurements with inventory data. Other approaches include comparisons of mobile source emissions based upon fuel consumption to estimates based on vehicle miles traveled, comparisons of emission trends estimated several years apart, comparisons with receptor-oriented model data, and inverse modeling using source-oriented models.²

Ambient measurements are often used because it is expected that a change in the emissions of a given pollutant will be reflected in a corresponding change in ambient concentrations, particularly when (unlike ozone and some forms of PM) there is little or no chemical transformation of the emissions. Instances in which temporal trends are not consistent between emissions and ambient concentration data can provide an indication of errors in one or both data sets or evidence that the link between emissions and ambient concentrations is not adequately understood. In some cases, ratios of ambient concentrations can be directly compared with ratios of emitted species. For example, the CO to NO_x and benzene to acetylene ratios have been used to evaluate mobile source emission inventories.^{2, 110, 111}

Fuel consumption data can also be used as the basis for comparing inventory data to related but independent data. Mobile source inventories are based upon vehicle miles traveled (VMT), with emissions estimated from measurements of emitted pollutant mass per unit distance. Where data are available that estimate pollutant mass emissions per unit fuel consumed, databases that report fuel consumption can be used as the basis for estimating mobile source emissions, and the results compared to the VMT-based emissions or ambient concentrations. This approach has been used to evaluate mobile source inventories in Nashville, TN and California.^{112, 113}

Receptor-oriented models estimate the contributions of specific source types to measured ambient pollutant concentrations by fitting measured concentrations of ambient species to a linear combination of source profiles. The contributions are in effect an inventory of emissions that are responsible for the ambient pollutant level at the location of the

receptor (an ambient monitoring site). The use of receptor models is only possible for pollutants for which there are differences in the relative mix of chemical species emitted by different source types. This technique has been used to evaluate inventories of PM_{2.5} and VOCs.^{7, 114}

Inverse modeling is an additional approach to evaluating emission inventory accuracy. Fundamentally, source-oriented air quality models use emission source strengths and other variables such as winds, solar insolation, and deposition rates as inputs into equations describing the physical and chemical processes taking place in the atmosphere to compute the ambient pollutant concentrations within the domain of interest. Inverse modeling involves reformulating the equations used in the model so that the emission source strengths are expressed in terms of the observed concentrations. For several reasons, the inverse modeling technique is usually applied to systems that are large in scale (global or continental) and that are characterized by relatively dispersed sources.^{66, 115, 116}

Top-down methods must be applied with caution to ensure that the comparisons are as appropriate as possible. Although the data used in top-down evaluations are (ideally) independently derived relative to the inventory data, top-down data are similarly subject to data uncertainty and limitations. Ambient data, for instance, include contributions from sources other than the source categories being evaluated, and care must be taken to verify that such contributions are minimal relative to the contributions of the categories of interest. The most effective applications of top-down evaluations are those that are combined with concurrent examination of the original bottom-up inventory data, so that the source of errors can be identified rather than simply stating that the inventory is in error.^{117, 118}

One of the most important insights from the literature on inventory uncertainty is that it is usually far more efficient and less resource-intensive to conduct uncertainty analyses at the time the data are developed and incorporated into the emission inventory methodology, rather than conducted after the fact.^{43, 104} After the fact, the original data

are often unavailable or are poorly documented, making it difficult or impossible to quantify uncertainty. Even when the original data are available and are adequately documented, the time required to conduct a retrospective evaluation can be prohibitive.

In addition to conducting a technical evaluation of data uncertainty, it is important to ensure that the process of incorporating data from the enormously wide range of data sources into a single inventory is made as seamless as possible. In the U.S., the process for developing the NEI is under evaluation with the purpose of making some fundamental changes to that process.^{119, 120} Other efforts are underway to develop regional inventories that draw upon the benefits of an open-source approach to allow users to make modifications to emission models and other algorithms, with the expectation that such an approach will result in a greater understanding of the inventory process and an increased potential for innovations in inventory data processing.¹²¹

Some of the possible changes are modification of the emission factor quality rating, ability to more routinely incorporate source test results into the emission factor database, and greater use of internet-based information exchange technologies such as eXtensible Markup Language (XML).^{2,122} Development and application of on-line tools and methods for exchanging, processing, and analyzing information are approaches that are currently widely used in most facets of business and throughout government, and will continue to be applied to inventory development and reporting.

CONCLUSIONS AND RECOMMENDATIONS

Although the development of emission inventories is often perceived as being a relatively straightforward process, it is in fact a complex and involved undertaking to collect, verify, and aggregate the wide range of data required to create the fundamental picture of the location and amount of pollutant emissions. The past (and ongoing) successes in improving air quality across North America are making it more difficult to maintain the gains already achieved and to continue the effort to protect the public from health problems caused or exacerbated by ambient air pollution, as the most readily controlled sources have been or are being controlled. This combination of process complexity and

increasing challenges make it imperative that the measurements, data processing, and modeling required to accurately estimate emissions of air pollutants take full advantage of the innovations that have been developed in recent years.

Numerous technologies and approaches have been demonstrated to be of considerable value to improving emission inventories, and are ready to be used to support recommendations made by several independent scientific panels to provide more accurate and timely information on where and when emissions occur that are of importance to maintaining and improving air quality. Although it is important to apply the technologies and methods now available, additional efforts are needed. In addition to further research, organizational efforts are also needed to make the collection and processing of inventory data more seamless.

The NARSTO Emission Inventory Assessment has identified eight recommendations to address the shortcomings of existing emission inventories.² They are:

1. *Reduce uncertainties associated with emissions from key undercharacterized sources:* Focus immediate measurement and development efforts on areas of greatest known uncertainty within current emission inventories. Systematically continue to improve emission inventories by applying sensitivity and uncertainty analyses and by comparing them to independent sources of measured data. Such comparisons will help identify subsequent improvement priorities.
2. *Improve speciation estimates:* Develop new and improve existing source speciation profiles and emission factors plus the related activity data needed to more accurately estimate speciated emissions for particulate matter and its precursors, volatile organic compounds, and air toxics.
3. *Improve existing and develop new emission inventory tools:* Continue the development of new and existing measurement and analysis technologies to enable expanded measurements of emissions and ambient concentrations. Apply these technologies in developing emission model and processor capabilities to allow models to more closely approximate actual emissions in time and space.

4. *Quantify and report uncertainty*: Develop guidance, measures, and techniques to improve uncertainty quantification, and include measures of uncertainty (including variability) as a standard part of reported emission inventory data.
5. *Increase inventory compatibility and comparability*: Define and implement standards for emission inventory structure, data documentation, and data reporting for North American emission inventories.
6. *Improve user accessibility*: Improve user accessibility to emission inventory data, documentation, and emission inventory models through the Internet or other electronic formats.
7. *Improve timeliness*: Create and support a process for preparing and reporting national emission inventory data on a yearly basis.
8. *Assess and improve emission projections*: Emission projection methodologies for all emission inventory sectors in North America should be evaluated to determine the accuracy of past projections and identify areas of improvement for future projections.

The priority of these recommendations will depend upon the particular situation facing each organization, whether at the federal, regional, state or provincial, or local levels. Nevertheless, each recommendation is applicable to all organizations.

Emission inventories face significant challenges in meeting the ever-increasing demands for timely and accurate information to address air quality management needs. Here many currently available technologies and approaches have been described that can improve emission inventories to enable them to meet those challenges. Applying these tools will require investments of time and money, but these investments will ensure that future air quality management decisions are based upon the best possible information. This, in turn, will lead to the development of air quality management strategies that are as effective as possible in terms of health, environmental, and economic measures.

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REFERENCES

1. Cass, G.R.; McRae, G.J. Minimizing the cost of air pollution control; *Environ. Sci. Technol.* 1981, 15, 748-757.
2. NARSTO. *Improving Emission Inventories for Effective Air Quality Management Across North America*; NARSTO: Oak Ridge, TN, 2005.
3. National Research Council. *Air Quality Management in the United States*; National Academy Press: Washington, DC, 2004.
4. McCabe, J.; Green, G. *Recommendations to the Clean Air Act Advisory Committee*; Air Quality Management Work Group; Washington, DC; January, 2005.
5. National Research Council. *Research Priorities for Airborne Particulate Matter: IV. Continuing Research Progress*; National Academy Press: Washington, DC, 2004.
6. National Research Council. *Modeling Mobile-Source Emissions*; National Academy Press: Washington, DC, 2000.
7. McMurry, P.H.; Shepherd, M.; Vickery, J., eds. *Particulate Matter Science for Policy Makers*. 2004, Cambridge University Press: New York, NY.
8. Mobley, J.D.; Cadle, S.H. Innovative methods for emission inventory development and evaluation: Workshop summary; *J. Air Waste Manage. Assoc.* 2004, 54(11), 1422-1439.
9. U.S. Environmental Protection Agency. *National Air Quality and Emissions Trends Report: 2003 Special Studies Edition*; EPA 454/R-03-005; Office of Air Quality Planning and Standards; Research Triangle Park, NC; September, 2003.
10. U.S. Environmental Protection Agency. Rule to reduce interstate transport of fine particulate matter and ozone (Clean air interstate rule); *Federal Register* 2005.
11. U.S. Environmental Protection Agency. Control of air pollution from new motor vehicles: Heavy-duty engine and vehicle standards and highway diesel fuel sulfur control requirements; Final rule; *Federal Register* 2001, 66(12), 5002.
12. U.S. Environmental Protection Agency. Control of emissions of air pollution from nonroad diesel engines and fuel; Final rule; *Federal Register* 2004, 69(124), 38958.
13. U.S. Environmental Protection Agency. *Clean Air Interstate Rule: Emissions Inventory Technical Support Document*; Office of Air Quality Planning and Standards; Research Triangle Park, NC; March 4, 2005.
14. Wark, K.; Warner, C.F. *Air Pollution, Its Origins and Control*; Harper and Row: New York, NY, 1976.
15. Stern, A.C.; Wohlers, H.C.; Boubel, R.W.; Lowry, W.P. *Fundamentals of Air Pollution*; Academic Press: New York, NY, 1973.
16. Werner, A.S.; Mobley, J.D. Emissions inventories then, now, and tomorrow; *EM* 2005, 41-44.

- 887 17. McCabe, L.C., ed. *Air Pollution: Proceedings of the United States Technical*
888 *Conference on Air Pollution*. 1952, McGraw-Hill: New York.
- 889 18. Rossano, A.T.; Schell, N.E. *Procedures for making an inventory of air pollution*
890 *emissions*. In *Golden Jubilee Meeting of the Air Pollution Control Association*.
891 June 3. Year. St. Louis, MO.
- 892 19. U.S. Environmental Protection Agency. *Compilation of Air Pollutant Emission*
893 *Factors: Volume I. Stationary Point and Area Sources, Fifth Edition*; AP-42;
894 Office of Air Quality Planning and Standards; Research Triangle Park, NC;
895 January, 1995.
- 896 20. Haagen-Smit, A.J. Air conservation; *Science* 1958, 128, 869-878.
- 897 21. U.S. Environmental Protection Agency. *Frequently Asked Questions on*
898 *MOBILE6*; EPA 420-B-03-013; Office of Transportation Air Quality; Ann Arbor,
899 MI; November, 2003.
- 900 22. U.S. Environmental Protection Agency. *Technical Guidance on the Use of*
901 *MOBILE6.2 for Emission Inventory Preparation*; EPA 420-R-04-013; Office of
902 Transportation Air Quality; Ann Arbor, MI; August, 2004.
- 903 23. U.S. Environmental Protection Agency. *User's Guide for the EPA Nonroad*
904 *Emissions Model Draft NONROAD 2002*; EPA 420-P-02-013; Office of
905 Transportation Air Quality; Ann Arbor, MI; December, 2002.
- 906 24. California Air Resources Board *EMFAC2001/EMFAC2002: Calculating*
907 *Emission Inventories for Vehicles in California: User's Guide*; California Air
908 Resources Board; Sacramento, CA, 2002.
- 909 25. Pierce, T.E.; Geron, C.; Pouliot, G.; Vukovich, J.; Kinnee, E. *Integration of the*
910 *Biogenic Emissions Inventory System (BEIS3) into the Community Multiscale Air*
911 *Quality (CMAQ) modeling system*. In *25th Agricultural and Forestry Meteorology*
912 *Conference*. May 20-24, 2002, Norfolk, VA.
- 913 26. Anderson, N.; Strader, R.; Davidson, C. Airborne reduced nitrogen: Ammonia
914 emissions from agriculture and other sources; *Environ. Int.* 2003, 29, 277-286.
- 915 27. U.S. Environmental Protection Agency. *MOBILE6: On-Road Motor Vehicle*
916 *Emissions Model, 5-Day Training Course*; 2001,
917 <http://www.epa.gov/otaq/models/mobile6/trng5day/sldday.pdf>, accessed
918 September 21, 2005.
- 919 28. Dykema, O.W.; Kemp, V.E. *Inventory of combustion-related emissions from*
920 *stationary sources*; EPA-600/7-76-012; Industrial Environmental Research
921 Laboratory; Research Triangle Park, NC; September, 1976.
- 922 29. Saeger, M. *The 1985 NAPAP Emissions Inventory (Version 2): Development of*
923 *the Annual Data and Modeler's Tapes*; EPA-600/7-89-012a; U.S. Environmental
924 Protection Agency; Research Triangle Park, NC, 1989.
- 925 30. U.S. Environmental Protection Agency. *2002 National Emission Inventory (NEI)*
926 *Preparation Plan, Final*; Office of Air Quality Planning and Standards; Research
927 Triangle Park, NC; August 10, 2004, 2004.
- 928 31. Allen, D.; Durenburger, C. *Accelerated science evaluation of ozone formation in*
929 *the Houston-Galveston area: Emission inventories*; 2003,
930 <http://www.utexas.edu/research/ceer/texaqsarchive/>, accessed February 5.

- 931 32. Kuhns, H.D.;Green, M.; Etyemezian, V. *Big Bend Regional Aerosol and Visibility*
932 *Observational (BRAVO) Study Emissions Inventory*; BRAVO Steering
933 Committee, Desert Research Institute; Reno, NV, 2001.
- 934 33. Kuhns, H.D.; Vukovich, J. *The emission inventories and SMOKE modeling efforts*
935 *used to support the BRAVO Study*. In *12th Annual U.S. EPA International*
936 *Emissions Inventory Conference*. April 29-May 1, 2003, San Diego, CA.
- 937 34. Eastern Research Group, *Border 2012 Emissions Inventory*. 2004, Western
938 Governors Association, EPA Region 9, EPA Office of International Activities.
- 939 35. U.S. Environmental Protection Agency. *Database of sources of environmental*
940 *releases of dioxin-like compounds in the United States*; EPA-600/C-01-012;
941 Office of Research and Development; Washington, DC; March, 2001.
- 942 36. Acosta-Ruiz, G.; Powers, B. *Preliminary Atmospheric Emissions Inventory of*
943 *Mercury for Mexico*. In *12th Annual U.S. EPA International Emissions Inventory*
944 *Conference*. April 29 - May 1, 2003, San Diego, CA.
- 945 37. Bond, T.C.;Streets, D.G.;Yarber, K.F.;Nelson, S.M.;Woo, J.-H.; Klimont, Z. A
946 technology-based global inventory of black and organic carbon emissions from
947 combustion; *J. Geophys. Res.* 2004, *109*, D14203.
- 948 38. National Research Council. *Rethinking the Ozone Problem in Urban and*
949 *Regional Air Pollution*; National Academy Press: Washington, DC, 1991.
- 950 39. Center for Environmental Modeling for Policy Development *SMOKE v2.1 User's*
951 *Manual*; 2005, <http://cf.unc.edu/cep/empd/products/smoke/version2.1/index.cfm>,
952 accessed September 21, 2005.
- 953 40. Adelman, Z.; Houyoux, M. *Processing the National Emissions Inventory 96*
954 *(NEI96) version 3.11 with SMOKE*. In *Emission Inventory Conference: One*
955 *Atmosphere, One Inventory, Many Challenges*. May, 2001, Denver, CO.
- 956 41. Hodan, W. *Updates and improvements to the SPECIATE data and program*. In
957 *12th Annual U.S. EPA International Emissions Inventory Conference*. April 29 -
958 May 1, 2003, San Diego, CA.
- 959 42. Placet, M.;Mann, C.O.;Gilbert, R.O.; Neifer, M.J. Emissions of ozone precursors
960 from stationary sources: A critical review; *Atmos. Environ.* 2000, *34*, 2183-2204.
- 961 43. Frey, H.C.; Zheng, J. Quantification of variability and uncertainty in utility NOx
962 emission inventories; *J. Air Waste Manage. Assoc.* 2002, *52*, 1083-1095.
- 963 44. Cullen, A.C.; Frey, H.C. *The use of probabilistic techniques in exposure*
964 *assessment: A handbook for dealing with variability and uncertainty in models*
965 *and inputs*; Plenum: New York, NY, 1999.
- 966 45. Morgan, M.G.; Henrion, M. *Uncertainty: A Guide to Dealing With Uncertainty*
967 *in Quantitative Risk and Policy Analysis*; Cambridge University Press: New York,
968 1990.
- 969 46. Childers, J.W.;Thompson, J., Edgar L.;Harris, D.B.;Kirchgessner, D.A.;Clayton,
970 M.;Natschke, D.F.; Phillips, W.J. Multi-pollutant concentration measurements
971 around a concentrated swine production facility using open-path FTIR
972 spectrometry; *Atmos. Environ.* 2001, *35*, 1923-1936.
- 973 47. Hashmonay, R.A.;Natschke, D.F.;Wagoner, K.;Harris, D.B.;Thompson, J., Edgar
974 L.; Yost, M.G. Field evaluation of a method for estimating gaseous fluxes from
975 area sources using open-path Fourier transform infrared; *Environ. Sci. Technol.*
976 2001, *35*(11), 2309-2313.

- 977 48. Yokelson, R.J.;Ward, D.E.;Susott, R.A.;Reardon, J.; Griffith, D.W.T. Emissions
978 from smoldering combustion of biomass measured by open-path Fourier
979 transform infrared spectroscopy; *J. Geophys. Res.* 1997, *102*, 18865-18877.
- 980 49. Haus, R.;Wilkinson, R.;Heland, J.; Schafer, K. Remote sensing of gas emissions
981 on natural gas flares; *Pure Appl. Opt.* 1998, *7*, 853-862.
- 982 50. Schäffer, K.;Emeis, S.;Hoffmann, H.; Jahn, C., *Emissions of Fuel Stations and*
983 *Tankers Determined by the Inverse Method*, in *Emissions of Air Pollutants --*
984 *Measurements, Calculations, and Uncertainties*, R. Fredrich and S. Reis, Editors.
985 2004, Springer: Berlin.
- 986 51. Schroter, M.;Obermeier, A.;Bruggemann, D.;Plehschmidt, M.; Klemm, O.
987 Remote monitoring of air-pollutant emissions from point sources by a mobile
988 lidar/sodar system; *J. Air Waste Manage. Assoc.* 2003, *53*, 716.
- 989 52. Bishop, G.A.;Starkey, J.R.;Ihlenfeldt, A.;Williams, W.J.; Stedman, D.H. IR long-
990 path photometry: A remote sensing tool for automobile emissions; *Anal. Chem.*
991 1989, *61*, 617A-676A.
- 992 53. Stedman, D.H. Automobile carbon monoxide emissions; *Environ. Sci. Technol.*
993 1989, *23*, 147-149.
- 994 54. Bishop, G.A.; Stedman, D.H. Measuring the emissions of passing cars; *Acc.*
995 *Chem. Research* 1996, *29*, 489-495.
- 996 55. Guenther, P.L.;Stedman, D.H.;Bishop, G.A.;Beaton, S.P.;Bean, J.H.; Quine, R.W.
997 A hydrocarbon detector for the remote sensing of vehicle exhaust emissions;
998 *Review of Scientific Instrumentation* 1995, *66*, 3024-3029.
- 999 56. Stephens, R.D.; Cadle, S.H. Remote sensing measurements of carbon monoxide
1000 emissions from on-road vehicles; *J. Air Waste Manage. Assoc.* 1991, *41*, 39-46.
- 1001 57. Jimenez, J.L.;Koplow, M.D.;Nelson, D.D.;Zahniser, M.S.; Schmidt, S.E.
1002 Characterization of on-road vehicle NO emissions by a TILDAS remote sensor; *J.*
1003 *Air Waste Manage. Assoc.* 1999, *49*, 463-470.
- 1004 58. Baum, M.M.;Kiyomiya, E.S.;Kumar, S.;Lappas, A.M.;Kapinus, V.A.; Lord III,
1005 H.C. Multicomponent remote sensing of vehicle exhaust by dispersive absorption
1006 spectroscopy: 2. Direct on-road ammonia measurements; *Environ. Sci. Technol.*
1007 2001, *35*, 3735-3741.
- 1008 59. Kuhns, H.D.;Mazzoleni, C.;Moosmuller, H.;Nikolic, D.;Keislar, R.E.;Barber,
1009 P.W.;Li, Z.;Etyemezian, V.; Watson, J.G. Remote sensing of PM, NO, CO, and
1010 HC emission factors for on-road gasoline and diesel engine vehicles in Las Vegas,
1011 NV; *Sci. Total Environ.* 2004, *322*, 123-137.
- 1012 60. Moosmuller, H.;Mazzoleni, C.;Barber, P.W.;Kuhns, H.D.;Keislar, R.E.; Watson,
1013 J.G. On-road measurement of automotive particle emissions by ultraviolet lidar
1014 and transmissometer instrument; *Environ. Sci. Technol.* 2003, *37*, 4971-4978.
- 1015 61. Baum, M.M.;Kiyomiya, E.S.;Kumar, S.;Lappas, A.M.; Lord III, H.C.
1016 Multicomponent sensing of vehicle exhaust by dispersive spectroscopy. 1. Effect
1017 of fuel type and catalyst performance; *Environ. Sci. Technol.* 2000, *34*, 2851-
1018 2858.
- 1019 62. Jiménez, J.L.;McManus, J.B.;Shorter, J.H.;Nelson, D.D.;Zahniser, M.S.;Koplow,
1020 M.D.;McRae, G.J.; Kolb, C.E. Cross road and mobile tunable infrared laser
1021 measurements of nitrous oxide emissions from motor vehicles; *Chemosphere*
1022 2000, *2*, 397-412.

- 1023 63. Soja, A.;Al-Saadi, J.;Pierce, B.;Szykman, J.;Williams, D.J.;Pace, T.;Kordzi, J.;
1024 Barnard, W.R. *Using satellite-based products to enhance existing area burned*
1025 *data*. In *14th International Emission Inventory Conference*. April 12-14, 2005,
1026 Las Vegas, NV.
- 1027 64. Palmer, P.I.;Jacob, D.J.;Fiore, A.M.;Martin, R.V.;Chance, K.; Kurosu, T.P.
1028 Mapping isoprene emissions over North America using formaldehyde column
1029 observations from space; *J. Geophys. Res.* 2003, *108*, 4180.
- 1030 65. Martin, R.V.;Jacob, D.J.;Chance, K.;Kurosu, T.P.;Palmer, P.I.; Evans, M.J.
1031 Global inventory of nitrogen oxide emissions constrained by space-based
1032 observations of NO₂ columns; *J. Geophys. Res.* 2003, *108*, 4537.
- 1033 66. Pétron, G.;Granier, C.;Khattatov, B.;Lamarque, J.-F.;Yudin, V.;Müller, J.-F.;
1034 Gille, J. Inverse modeling of carbon monoxide surface emissions using CMDL
1035 network observations; *J. Geophys. Res.* 2002, *107*, 4761.
- 1036 67. Melamed, M.L.;Solomon, S.;Daniel, J.S.;Langford, A.O.;Portmann,
1037 R.W.;Ryerson, T.B.;Nicks Jr., D.K.; McKeen, S.A. Measuring reactive nitrogen
1038 emissions from point sources using visible spectroscopy from aircraft; *J Environ*
1039 *Monitor* 2002, *5*, 29-34.
- 1040 68. Stearns, J.R.;Zahniser, M.S.;Kolb, C.E.; Sanford, B.P. Airborne infrared
1041 observations and analyses of a large forest fire; *Appl Optics* 1986, *25*, 2554-2562.
- 1042 69. Sawyer, R.;Harley, R.A.;Cadle, S.H.;Norbeck, J.;Slott, R.; Bravo, H. Mobile
1043 sources critical review: 1998 NARSTO Assessment; *Atmos. Environ.* 2000, *34*,
1044 2161-2181.
- 1045 70. U.S. Environmental Protection Agency. *A Roadmap to MOVES2004*; EPA-420-S-
1046 05-002; Office of Transportation Air Quality; Ann Arbor, MI; March, 2005.
- 1047 71. Bishop, G.A.; Stedman, D.H. On-road carbon monoxide emission measurement
1048 comparisons for the 1988-1989 Colorado oxy-fuels program; *Environ. Sci.*
1049 *Technol.* 1990, *24*, 843-847.
- 1050 72. National Research Council. *Evaluating Vehicle Emissions Inspection and*
1051 *Maintenance Programs*; National Academy Press: Washington, DC, 2001.
- 1052 73. Stephens, R.D. Remote sensing data and a potential model of vehicle exhaust
1053 emissions; *J. Air Waste Manage. Assoc.* 1994, *44*, 1284-1292.
- 1054 74. Mazzoleni, C.;Moosmüller, H.;Kuhns, H.D.;Keislar, R.E.;Barber, P.W.;Nikolic,
1055 D.;Nussbaum, N.J.; Watson, J.G. Correlation between automotive CO, HC, NO,
1056 and PM emission factors from on-road remote sensing: implications for inspection
1057 and maintenance programs; *Transport Res* 2004, *D9*, 477-496.
- 1058 75. California Bureau of Automotive Repair *Device High Emitter Identification with*
1059 *Confirmatory Roadside Inspection, Final Report*; Report 2001-06; Engineering
1060 and Research Branch; Sacramento, CA, 2001.
- 1061 76. Bishop, G.A.;Morris, J.A.;Stedman, D.H.;Cohen, L.H.;Countess, R.G.;Maly, P.;
1062 Scherer, S. The effects of altitude on heavy duty diesel truck on-road emissions;
1063 *Environ. Sci. Technol.* 2001, *35*, 1574-1578.
- 1064 77. Jiménez, J.L.;McRae, G.J.;Nelson, D.D.;Zahniser, M.S.; Kolb, C.E. Remote
1065 sensing of NO and NO₂ emissions from heavy-duty diesel trucks using tunable
1066 diode lasers; *Environ. Sci. Technol.* 2000, *34*, 2380-2387.

- 1067 78. Bishop, G.A.;Morris, J.A.; Stedman, D.H. Snowmobile contributions to mobile
1068 source emissions in Yellowstone National Park; *Environ. Sci. Technol.* 2001, 35,
1069 2874-2881.
- 1070 79. Wenzel, T.;Singer, B.C.; Slott, R.S. Some issues in the statistical analysis of
1071 vehicle emissions; *J Transport Stat* 2000, 3, 1-4.
- 1072 80. Pierson, W.; Brachaczek, W. Particulate matter associated with vehicles on the
1073 road; *Aerosol Sci. Technol.* 1983, 2, 1-40.
- 1074 81. Gertler, A.;Sagebiel, J.;Wittorff, D.;Pierson, W.;Dipple, W.;Freeman, W.; Sheetz,
1075 L. *Vehicle Emissions in Five Urban Tunnels*; Report for Coordinating Research
1076 Council; Desert Research Institute; Reno, NV, 1997.
- 1077 82. Pierson, W.;Gertler, A.; Bradow, R. Comparison of the SCAQS tunnel study with
1078 other on-road vehicle emission data; *J. Air Waste Manage. Assoc.* 1990, 40, 1495-
1079 1504.
- 1080 83. Graham, L.A.;Gray, C.;Rogak, S.; Brakel, T. *Pacific 2001: Cassiar Tunnel study-
1081 Particulate matter emissions measurements. In the NARSTO Emission Inventory
1082 Workshop, Innovative Methods for Emissions Inventory Development and
1083 Evaluation.* 2003, Austin, TX.
- 1084 84. Kolb, C.E.;Herndon, S.C.;McManus, J.B.;Shorter, J.H.;Zahniser, M.S.;Nelson,
1085 D.D.;Jayne, J.T.;Canagaratna, M.R.; Worsnop, D.R. Mobile laboratory with rapid
1086 response instruments for real-time measurements of urban and regional trace gas
1087 and particulate distributions and emission source characteristic; *Environ. Sci.
1088 Technol.* 2004, 38, 5694-5703.
- 1089 85. Seakins, P.W.;Lansley, D.L.;Hodgson, A.;Huntley, N.; Pope, F. New directions:
1090 Mobile laboratory reveals new issues in urban air quality; *Atmos. Environ.* 2002,
1091 36, 1247-1248.
- 1092 86. Bukowiecki, N.;Dommen, J.;Prévôt, A.S.H.;Richter, R.;Weingartner, E.;
1093 Baltensperger, U. A mobile pollutant measurement laboratory - measuring gas
1094 phase and aerosol ambient concentrations with high spatial and temporal
1095 resolution; *Atmos. Environ.* 2002, 36, 5569-5579.
- 1096 87. Kittleson, D.B.;Watts, J., W.F.; Johnson, J.P. Nanoparticle emissions on
1097 Minnesota highways; *Atmos. Environ.* 2004, 38, 9-19.
- 1098 88. Gouriou, F.;Morin, J.P.; Weill, M.-E. On-road measurements of particle number
1099 concentrations and size distributions in urban and tunnel environments; *Atmos.
1100 Environ.* 2004, 38, 2831-2840.
- 1101 89. Weijers, E.P.;Khlystov, A.Y.;Kos, G.P.A.; Erisman, J.W. Variability of
1102 particulate matter concentrations along roads and motorways determined by a
1103 moving measurement unit; *Atmos. Environ.* 2004, 38, 2993-3002.
- 1104 90. Kittleson, D.B.;Johnson, J.;Watts, W.;Wei, Q.;Drayton, M.;Paulsen, D.;
1105 Bukowiecki, N. *Diesel aerosol sampling in the atmosphere*; SAE Technical Paper
1106 Series 2000-01-2212, 2000.
- 1107 91. Shorter, J.H.;Herndon, S.C.;Zahniser, M.S.;Nelson, D.D.;Jayne, J.T.; Kolb, C.E.
1108 *Characterization of heavy-duty vehicle exhaust in dense urban environments. In
1109 10th International Symposium on Transport and Air Pollution.* Year. Boulder,
1110 CO.

- 1111 92. Vogt, R.;Scheer, V.;Casati, R.; Benter, T. On-road measurement of particle
1112 emissions in the exhaust plume of a diesel passenger car; *Environ. Sci. Technol.*
1113 2003, 37, 4070-4076.
- 1114 93. Canagaratna, M.R., Jayne, J.T., Ghertner, A., Herndon, S.C., Shi, Q., Jiménez,
1115 J.L., Silva, P., Williams, P., Lanni, T., Drewnick, F., et al. Chase studies of
1116 particulate emissions from in-use New York City vehicles; *Aerosol Sci. Technol.*
1117 2004, 38, 555-573.
- 1118 94. U.S. Environmental Protection Agency., *Real-time on-road vehicle exhaust gas*
1119 *modular flowmeter and emissions reporting system*, U.S. Patent Office, Editor.
1120 2003: United States.
- 1121 95. Baldauf, R.W.;Somers, J.S.;Tierney, G.;Fulper, C.R.;Warila, J.;Gabele, P.;Bailey,
1122 B.;Cadle, S.H.; Lawson, D.R. *Assessing particulate matter emissions from light-*
1123 *duty, gasoline powered motor vehicles*. In *13th Annual Emission Inventory*
1124 *Conference*. June, 2004, Clearwater, FL.
- 1125 96. Banet, M. *Identification of excessive emissions system failure rates in high-*
1126 *mileage fleet vehicles based on Networkcar's continuous on-board emissions*
1127 *monitoring system*. In *NARSTO Conference on Innovative Methods for Emission*
1128 *Inventory Development and Evaluation*. October, 2003, Austin, TX.
- 1129 97. Vogt, M.C.;Shoemaker, E.L.; Fraioli, A.V., *Electrocatalytic Cermet Gas*
1130 *Detector/Sensor*, U.S.P.a.T. Office, Editor. 1995.
- 1131 98. Gschwandtner, G. Trends and uncertainties in anthropogenic VOC and NO_x
1132 emissions; *Journal of Water, Air and Soil Pollution* 1993, 67, 39-46.
- 1133 99. Chang, W.;Cardelino, C.; Chang, M. The use of survey data to investigate ozone
1134 sensitivity to point sources; *Atmos. Environ.* 1996, 30(23), 4095-4099.
- 1135 100. Lee, D.S.;Kholler, I.;Grobler, E.;Rohrer, F.;Sausen, R.;Klenner, L.;Olivier,
1136 J.G.J.;Dentener, F.J.; Bouwman, A.F. Estimation of global NO_x emissions and
1137 their uncertainties; *Atmos. Environ.* 1997, 31, 1735-1749.
- 1138 101. van Amstel, A.;Olivier, J.G.J.; Ruysenaars, P.G. *Monitoring of greenhouse gases*
1139 *in the Netherlands: uncertainty and priorities for improvement*. In *Proceedings of*
1140 *a National Workshop*. Year. Bilthoven, The Netherlands: RIVM.
- 1141 102. El-Fadel, M.;Zeinati, M.;Ghaddar, N.; Mezher, T. Uncertainty in estimating and
1142 mitigating industrial related GHG emissions; *Energ Policy* 2001, 29, 1031-1043.
- 1143 103. Hanna, S.R.; Wilkinson, J. *Analytical estimation of uncertainties in biogenic*
1144 *emissions calculated by BEIS3 due to uncertainties in model inputs and*
1145 *parameters*. In *13th Annual Emission Inventory Conference*. 2004, Clearwater,
1146 FL.
- 1147 104. Frey, H.C.; Zheng, J. Probabilistic analysis of driving cycle-based highway
1148 vehicle emission factors; *Environ. Sci. Technol.* 2002, 36(23), 5184-5191.
- 1149 105. Hanna, S.R.;Lu, Z.;Chang, J.C.;Fernau, M.; Hansen, D.A. Monte Carlo estimates
1150 of uncertainties in predictions by a photochemical grid model (UAM-IV) due to
1151 uncertainties in input variables; *Atmos. Environ.* 1998, 32(21), 3619-3628.
- 1152 106. Bergin, M.;Noblet, G.;Petrini, K.;Dhieux, J.;Milford, J.; Harley, R.A. Formal
1153 uncertainty analysis of a Lagrangian photochemical air pollution model; *Environ.*
1154 *Sci. Technol.* 1999, 33, 1116-1126.
- 1155 107. Hanna, S.R.;Lu, Z.;Frey, H.C.;Wheeler, N.;Vukovich, J.;Arunachalam, S.;Fernau,
1156 M.; Hansen, D.A. Uncertainties in predicted ozone concentrations due to input

- 1157 uncertainties for the UAM-V photochemical grid model applied to the July 1995
 1158 OTAG domain; *Atmos. Environ.* 2001, 35(5), 891-903.
- 1159 108. Moore, G.; Londergan, R. Sampled Monte Carlo uncertainty analysis for
 1160 photochemical grid models; *Atmos. Environ.* 2001, 35, 4863-4876.
- 1161 109. Abdel-Aziz, A.; Frey, H.C. Propagation of Uncertainty in Hourly Utility NO_x
 1162 Emissions through a Photochemical Grid Air Quality Model: A Case Study for
 1163 the Charlotte, NC, Modeling Domain; *Environ. Sci. Technol.* 2004, 38, 2153-
 1164 2160.
- 1165 110. Fujita, E.M.; Croes, B.E.; Bennett, C.L.; Lawson, D.R.; Lurmann, F.W.; Main, H.H.
 1166 Comparison of emission inventory and ambient concentration ratios of CO,
 1167 NMOG, and NO_x in California's South Coast Air Basin; *J. Air Waste Manage.*
 1168 *Assoc.* 1992, 42, 264-276.
- 1169 111. Parrish, D.D.; Trainer, M.; Hereid, D.; Williams, E.J.; Olszyna, K.J.; Harley,
 1170 R.A.; Meagher, J.F.; Fehsenfeld, F.C. Decadal change in carbon monoxide to
 1171 nitrogen oxide ratio in U.S. vehicular emissions; *J. Geophys. Res.* 2002, D12,
 1172 4140.
- 1173 112. Harley, R.A.; McKeen, S.A.; Pearson, J.; Rodgers, M.O.; Lonneman, W.A.
 1174 Analysis of motor vehicle emissions during the Nashville/Middle Tennessee
 1175 ozone study; *J. Geophys. Res.* 2001, 106(D4), 3559-3567.
- 1176 113. Singer, B.C.; Harley, R.A. A fuel-based inventory of motor vehicle exhaust
 1177 emissions in the Los Angeles area during summer 1997; *Atmos. Environ.* 2000,
 1178 34, 1783-1795.
- 1179 114. Watson, J.G.; Chow, J.C.; Fujita, E.M. Review of volatile organic compound
 1180 source apportionment by chemical mass balance; *Atmos. Environ.* 2001, 1567-
 1181 1584.
- 1182 115. Park, R.J.; Jacob, D.J.; Chin, M.; Martin, R.V. Sources of carbonaceous aerosols
 1183 over the United States and implications for natural visibility; *J. Geophys. Res.*
 1184 2003, 108, 4355.
- 1185 116. Gilliland, A.B.; Dennis, R.L.; Roselle, S.J.; Pierce, T.E. Seasonal NH₃ emission
 1186 estimates for the Eastern United States using ammonium wet concentrations
 1187 and an inverse modeling method; *J. Geophys. Res.* 2003, 108, 4477.
- 1188 117. Gilliland, A.B.; Dennis, R.L.; Roselle, S.J.; Pinder, R.W. *Inverse model estimation*
 1189 *of seasonal NH₃ emissions.* In *2005 AAAR PM Supersites Conference: Particulate*
 1190 *Matter Supersites Program and Related Studies.* February, 2005, Atlanta, GA.
- 1191 118. Pinder, R.W.; Adams, P.J.; Gilliland, A.B. *Improvements to regional air quality*
 1192 *modeling from recent advances in ammonia emission inventory development.* In
 1193 *2005 AAAR PM Supersites Conference: Particulate Matter Supersites Program*
 1194 *and Related Studies.* February, 2005, Atlanta, GA.
- 1195 119. Solomon, D.; Dombrowski, S.; Harwell, L.; Tillerson, C.; Wagoner, R. *The Rapid*
 1196 *Inventory Development Pilot.* In *14th International Emission Inventory*
 1197 *Conference.* April 12-14, 2005, Las Vegas, NV.
- 1198 120. Solomon, D. *Reinventing the NEI: A Status Report.* In *14th International*
 1199 *Emission Inventory Conference.* April 12-14, 2005, Las Vegas, NV.
- 1200 121. Janssen, M. *Transparent, Comprehensive, and Ready for Modeling - Building*
 1201 *Regional Inventories in the 21st Century.* In *14th International Emission*
 1202 *Inventory Conference.* April 12-14, 2005, Las Vegas, NV.

1203 122. Lane, B.E.;Ramachandran, V.;Lettich, R.;Minnich, K.;Sarode, A.; Garofalo, B.
1204 *Presentation of the AES*Online and AES*XML Emission Inventory Application.*
1205 *In 14th International Emission Inventory Conference.* April 12-14. Year. Las
1206 Vegas, NV.
1207

1208 **TABLE**1209 **Table 1.** Estimated relative confidence levels of national emission inventory data for selected pollutants.²1210 Confidence levels for SO₂, NO_x, and VOC are from the NARSTO PM Assessment.⁷

Pollutant	Source	Estimated Confidence Levels in Overall Inventory		
		Canada	U.S.A	Mexico
SO ₂	Utilities	high	high	high
	Other point sources	medium	medium	low-medium
	On-road mobile	medium	medium	low
	Non-road mobile	Low-medium	medium	low
	Area sources	low	low	low
	Biogenic source	low	low	low
	Other man-made sources (non-combustion)	low	low	low
NO _x	Utilities	medium-high	high	medium
	Other point sources	medium	medium	medium
	On-road mobile	high	high	medium
	Non-road mobile	medium-high	medium-high	low
	Area sources	low	low	low
	Biogenic source	low	low	low
	Other man-made sources (non-combustion)	medium	medium	low
VOC	Utilities	medium-high	medium-high	medium
	Other point sources	low-medium	low-medium	medium
	On-road mobile	medium	medium-high	low
	Non-road mobile	medium	Medium	low
	Area sources	low	Low	low
	Biogenic source	low	low	low
	Other man-made sources (non-combustion)	medium	medium	low
HAP	Utilities	medium	medium	medium
	Other point sources	low-medium	low-medium	low
	On-road mobile	medium	medium	low
	Non-road mobile	low-medium	low-medium	low
	Area sources	low	low	low
	Biogenic source	low	low	low
	Other man-made sources (non-combustion)	low	low	low

1211

FIGURE CAPTIONS

Figure 1. Estimates (2001) and projections (2020) of annual U.S. emissions of NO_x (left) and SO₂ (right).⁸

Figure 2. Schematic of the structure of an emission inventory that uses emission and activity factors and emission models.²⁶

Figure 3. Schematic of data flow in SMOKE emission processor. A linear processor would apply each program consecutively rather than in parallel.³⁸

Emission inventories are the starting point for managing air quality. Shortcomings in data and methods used to develop current emission inventories can lead to potentially ineffective air quality management strategies. By understanding these shortcomings (and emission inventory strengths), air quality managers can identify what new technologies can be applied and what additional data are most likely to provide the greatest improvement in airshed characterization. The recommendations provide a guide for what improvements are most important and most likely to result in improved air quality management capabilities.

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